

TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED / ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		ATTORNEY'S DOCKET NUMBER P67430US0
		US APPLICATION NO. (if known, see 37 CFR 1.5) 10/019465
INTERNATIONAL APPLICATION NO. ✓ PCT/DK00/00361	INTERNATIONAL FILING DATE 3 July 2000	PRIORITY DATE CLAIMED 2 July 1999
TITLE OF INVENTION AN ACCESS MEMBER AND A SYSTEM FOR CATHETERIZATION OF THE URINARY BLADDER THROUGH AN 'ARTIFICIAL OR A NATURAL CANAL IN A USER, AND A METHOD OF REPLACING SUCH AN ACCESS MEMBER		
APPLICANT(S) FOR DO/EO/US Morten Bay ALEXANDERSEN -and- Suzanne Eis BENZON		

Applicant herein submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information.

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for Internatl. Preliminary Examination was made by the 19th month from earliest claimed priority date.
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US)
6. ☐ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ A translation of the annexes to the Internatl. Preliminary Examination report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11. to 16. below concern other document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A **FIRST** preliminary amendment.
☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
14. ☐ A substitute specification.
15. ☐ A change of power of attorney and/or address letter.
16. ☒ Other items or information:
 - International Search Report – EPO
 - PCT Request Form
 - PCT/IB/304 Form
 - PCT/IB/308 Form
 - First Page of Publication
 - International Preliminary Examination Report – with Annexes

US APPLICATION NO. (if known, see 37 CFR 1.5) 10/019465		INTERNATIONAL APPLICATION NO. PCT/DK00/00361		ATTORNEY'S DOCKET NUMBER P67430US0	
<p>17. <input checked="" type="checkbox"/> The following fees are submitted:</p> <p>Basic National Fee (37 CFR 1.492(a)(1)-(5)):</p> <p>Internatl. prelim. examination fee paid to USPTO (37 CFR 1.492 (a) (1)) . . \$710.00</p> <p>No international preliminary examination fee paid to USPTO (37 CFR 1.492 (a) (2)) but international search fee paid to USPTO (37 CFR 1.445(a)(2)) . . \$740.00</p> <p>Neither international preliminary examination fee (37 CFR 1.492 (a) (3)) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO) \$1040.00</p> <p>International preliminary examination fee paid to USPTO (37 CFR 1.492 (a) (4)) and all claims satisfied provisions of PCT Article 33(2)-(4) \$100.00</p> <p>Search Report prepared by the EPO or JPO (37 CFR 1.492 (a) (5)) \$890.00</p> <p style="text-align: center;">ENTER APPROPRIATE BASIC FEE AMOUNT =</p>				CALCULATIONS	PTO USE ONLY
<p>Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input checked="" type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).</p>				\$ 130.00	
Claims	Number Filed	Number Extra	Rate		
Total Claims	18 - 20 =	-0-	x \$18.00	\$	
Independent Claims	1 - 3 =	-0-	x \$84.00	\$	
Multiple Dependent Claim(s) (if applicable)			+ \$280.00	\$	
TOTAL OF ABOVE CALCULATIONS =				\$ 1020.00	
<p>Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 CFR 1.9, 1.27, 1.28).</p>				\$	
SUBTOTAL =				\$ 1020.00	
<p>Processing fee of \$130 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f))</p>				\$	
TOTAL NATIONAL FEE =				\$ 1020.00	
<p>Fee of \$40.00 for recording the enclosed assignment (37 CFR 1.21(h)). Assignment must be accompanied by appropriate cover sheet (37 CFR 3.28, 3.31).</p>				\$	
TOTAL FEES ENCLOSED =				\$ 1020.00	
				Amt. to be refunded:	\$
				Amt. charged:	\$
<p>a. <input checked="" type="checkbox"/> A check in the amount of \$ <u>1020.00</u> to cover the above fees is enclosed.</p> <p>b. <input type="checkbox"/> Please charge my Deposit Account No. <u>06-1358</u> in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.</p> <p>c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge my account any additional fees set forth in §1.492 during the pendency of this application, or credit any overpayment to Deposit Account No. <u>06-1358</u>. A duplicate copy of this sheet is enclosed.</p>					
<p>SEND ALL CORRESPONDENCE TO:</p> <div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>JACOBSON HOLMAN PLLC 400 7th Street, N.W., Suite 600 Washington, DC 20004 202-638-6666 CUSTOMER NUMBER: 00136</p> </div> <div style="width: 45%; text-align: right;"> <p>By <i>Jonathan I. Schenck</i> Reg. No. 29,151 for Harvey B. Jacobson, Jr. Reg. No. 20,851</p> </div> </div>					

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Morten Bay ALEXANDERSEN et al

Serial No.: New

Filing Date: December 31, 2001

For: AN ACCESS MEMBER AND A SYSTEM FOR CATHETERIZATION
OF THE URINARY BLADDER THROUGH AN ARTIFICIAL OR A
NATURAL CANAL IN A USER, AND A METHOD OF REPLACING
SUCH AN ACCESS MEMBER

PRELIMINARY AMENDMENT

Assistant Commissioner of Patents
Washington, D.C. 20231

Sir:

Prior to initial examination, please amend the above-
identified application as follows:

IN THE SPECIFICATION

Please insert the following sentence on line 1, immediately
following the title:

--This is a nationalization of PCT/DK00/00361, filed July 3,
2000 and published in English.--

IN THE CLAIMS

Please amend claims 4-6, 8-13 and 16-17 as follows:

4. (amended) An access member according to claim 1, characterized in that at least one part of the wall or walls of the access member comprises a net material of eg. metal.

5. (amended) An access member according to claim 1, characterized by comprising one wall forming a substantially hose-shaped access member.

6. (amended) An access member according to claim 1, characterized by comprising at least two walls which are formed by sheets of material having substantially larger dimensions in the longitudinal direction than in the transverse direction and being joined at the respective longitudinally extending edges.

8. (amended) An access member according to claim 6, characterized in that said sheets have different thicknesses.

9. (amended) An access member according to claim 6, characterized in that said sheets have different degrees of flexibility.

10. (amended) An access member according to claim 6, characterized in that at least one blind hold is provided in at least one of said sheets.

11. (amended) An access member according to claim 6, in which there are at least three sheets and two cavities, characterized in that one of said cavities is closed at a distance from the outer end of the access member.

12. (amended) An access member according to claim 1, characterized in that the inner end of the access member is designed as a cap having a number of openings.

13. (amended) An access member according to claim 1, characterized by comprising means for securing the outer end of the access member to the abdominal skin surface.

16. (amended) An access member according to claim 1, characterized in that a plug member is provided for insertion into the outer end of said at least one through-going cavity.

17. (amended) A system for catheterization of the urinary bladder through an artificial or a natural canal in a user, comprising a catheter adapted to be inserted through the canal, and an access member according to claim 1.

REMARKS

The foregoing Preliminary Amendment is requested in order to delete the multiple dependent claims and avoid paying the multiple dependent claims fee.

Attached hereto is a marked-up version of the changes made to the specification and claims by the current amendment. The attached page is captioned "VERSION WITH MARKINGS TO SHOW CHANGES MADE."

Early action on the merits is respectfully requested.

Respectfully submitted,

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Atty. Docket: P67430US0
Date: December 31, 2001
HBJ:jrc

VERSION WITH MARKINGS TO SHOW CHANGES MADE

IN THE CLAIMS

4. (amended) An access member according to claim 1 ~~any of claims 1 to 3~~, characterized in that at least one part of the wall or walls of the access member comprises a net material of eg. metal.
5. (amended) An access member according to claim 1 ~~any of claims 1 to 4~~, characterized by comprising one wall forming a substantially hose-shaped access member.
6. (amended) An access member according to claim 1 ~~any of claims 1 to 4~~, characterized by comprising at least two walls which are formed by sheets of material having substantially larger dimensions in the longitudinal direction than in the transverse direction and being joined at the respective longitudinally extending edges.
8. (amended) An access member according to claim 6 ~~to 7~~, characterized in that said sheets have different thicknesses.
9. (amended) An access member according to claim 6 ~~any of claims 6 to 8~~, characterized in that said sheets have different degrees of flexibility.
10. (amended) An access member according to claim 6 ~~any of claims 6 to 9~~, characterized in that at least one blind hold is provided in at least one of said sheets.

11. (amended) An access member according to claim 6 ~~any of claims 6 to 9~~, in which there are at least three sheets and two cavities, characterized in that one of said cavities is closed at a distance from the outer end of the access member.

12. (amended) An access member according to claim 1 ~~any of the preceding claims~~, characterized in that the inner end of the access member is designed as a cap having a number of openings.

13. (amended) An access member according to claim 1 ~~any of the preceding claims~~, characterized by comprising means for securing the outer end of the access member to the abdominal skin surface.

16. (amended) An access member according to claim 1 ~~any of the preceding claims~~, characterized in that a plug member is provided for insertion into the outer end of said at least one through-going cavity.

17. (amended) A system for catheterization of the urinary bladder through an artificial or a natural canal in a user, comprising a catheter adapted to be inserted through the canal, and an access member according to claim 1 ~~any of claims 1 to 16~~.

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An access member and a system for catheterization of the urinary bladder through an artificial or a natural canal in a user, and a method of replacing such an access member.

5

BACKGROUND OF THE INVENTION.

The present invention relates to an access member and a system for catheterization of the urinary bladder through an artificial or a natural canal in a user. The invention furthermore relates to a method of replacing such an access member

Catheterization may typically be necessary in the case of postoperative urine retention of newly operated patients in a hospital. Another typical use is with patients suffering from severe cases of urinary incontinence as for disabled individuals like para- or tetraplegics who frequently have no control permitting voluntary urination.

Traditionally, such catheterization is carried out by inserting a catheter through the urethra of the patient. The catheter may be left in place for permanent catheterization during several hours or days, which is typically the case in elderly and infirm patients, or be retracted after emptying of the bladder, ie. so-called intermittent catheterization (IC).

Access to the urinary bladder may likewise be desirable in order to introduce eg. pharmaceuticals into the bladder or in order to wash or rinse the bladder.

Intermittent urethral catheterization performed with intervals of eg. 3 to 6 hours reduces the risk of infection of urethra and the bladder significantly as compared to permanent catheterization and has for many users become increasingly common also in daily life

situations outside the clinical environment of a hospital, whereby a significantly improved quality of life has been obtained for this group of patients.

However, intermittent catheterization requires a certain degree of dexterity and mobility which implies that self-catheterization is not always possible, especially in women where the urethral orifice may be difficult to locate.

During recent years, suprapubic catheterization (SPC) has been introduced as an alternative to urethral catheterization. In suprapubic catheterization, a canal is made from the surface skin of the abdominal wall of a user into the bladder under local or general anaesthesia and by means of a pointed hollow introducer or trocar. After penetration of the trocar into the bladder, a catheter is inserted through the canal thus provided, the inner end of said catheter being retained in the bladder by means of eg. an inflatable balloon abutting the inner wall of the bladder after retraction of the trocar. Although many of the disadvantages connected with urethral catheterization, such as eg. urethral cleavage and urethritis, may be overcome by this technique, infection risk is still high as suprapubic catheterization is typically performed as permanent catheterization due to the fact that the canal may close during replacement of the catheter. Furthermore, the fact that the end of the catheter protrudes well into the bladder when using a balloon, which is necessarily placed at a distance from the end in order to allow in-flow of urine, means that the bladder wall may be injured, the more so as the bladder wall often assumes an at least partially collapsed position in which it rests on the end of the catheter.

GB patent No. 2 275 420 discloses a system for suprapubic catheterization of the bladder permitting

intermittent catheterization by means of an accessor or sealing member permanently lodged in the artificial canal. The accessor comprises an outer shell formed by two elongate leaves of a bendable plastics material which are hinged together along one edge and having flanges at one end for securing the accessor to the skin surface. A sealing means in the form of a balloon assembly keeps the canal formed in the accessor closed between emptyings but allows insertion of the catheter. Due to the size and material of the accessor, this system may cause discomfort to the user.

Another alternative is provided by the so-called Mitrofanoff principle, by which a suprapubic canal is surgically made by removing parts of a body section, such as the appendix, another part of the intestinal system, eg. a section of the ileum, or any other suitable tubular body tissue, and subsequently attaching one end of the section to the abdominal skin surface whereas the other end penetrates the bladder wall and possibly protrudes into the bladder, the part being attached to the bladder wall at the point of penetration. Obviously, this technique requires surgery under general anaesthesia and implies a loss of bowel or other tissue as well as stitches in the bladder wall.

US 5,704,353 discloses a catheter for temporary placement in the female urethra. The catheter comprises a shaft which in one end has a sealing portion and in the other end a cap. In the lumen of the shaft a one-way valve is enclosed, urine being drained upon activation of the valve by means of a spike. As the length of the shaft has to be adapted to the individual length of the user's urethra and due to the rather elaborate design, this device is expensive and complicated in manufacture. Furthermore, the presence

of the sealing portion, which is designed as a mushroom-shaped crown and which in the position of use rests against the inner surface of the bladder, may cause discomfort to the user.

5

SUMMARY OF THE INVENTION.

It is an object of the present invention to provide an access member for use in catheterization of the urinary bladder, which is comfortable to wear and
10 which at the same time provides for an appropriate security against leakage.

It is a further object to provide an access member, by which intermittent catheterization may be performed by a larger group of users and which
15 alleviates the problems encountered in the prior art.

These and other objects are met by an access member adapted to be, in a position of use, accommodated in an artificial or a natural canal in a user, said access member having an outer end and an
20 inner end defining a predetermined length and extending, in the position of use, from the outside of the body of the user through said canal and into the urinary bladder, and comprising at least one wall defining at least one cavity extending substantially
25 throughout said predetermined length, said at least one cavity being intended for intermittently receiving a catheter, said access member being characterized in that said at least one wall of the access member has such a degree of flexibility that said at least one
30 cavity is kept in a substantially closed position by the mutual contact of parts of said at least one wall, but allows for intermittent insertion of a catheter.

The flexibility of the wall or walls of the access member entail that the access member itself is
35 able to provide for the necessary sealing properties,

4a

as the access member will inherently have the effect of
an automatic non-return valve. In case the access
member is exposed to forces in the radial or
longitudinal directions, the wall of the access member
5 is pressed against itself or, alternatively, the walls
are pressed against each other, thus closing the
through-going cavity of the access member between
catheterizations, either by a collapse in the radial
direction and/or by a bend at the entrance into the
10 bladder. At the inner end of the access member, the
cavity is kept closed eg. by contraction of the
detrusor and possibly by the pressure exerted by the
urine collected in the bladder. By integrating the
sealing properties in the access member, it is possible
15 to make the access member according to the invention

very comfortable to wear.

During use in connection with suprapubic catheterization, in which the body canal is an artificial canal extending from the user's abdominal wall, contraction of the abdominal muscles keeps the part of the through-going cavity of the access member, which is situated in the region of the abdominal wall, closed, so that urine may not penetrate to the outside and consequently that eg. water may not seep into the bladder when the user is washing or bathing. Nevertheless, intermittent catheterization may be carried out without difficulty by inserting the catheter through the passage provided by the cavity or cavities of the access member.

In relation to the Mitrofanoff principle, the access member according to the invention does not necessarily require surgery under general anaesthesia or any loss of body tissue. By this design an access member is provided which makes intermittent catheterization a feasible and/or attractive alternative to a large number of users which hitherto have been forced to use permanent catheterization. As a consequence, it is possible to reduce the risk of infection in this group of users.

The wall or walls of the access member may comprise a foil or film material, or a foam or a gel. It is likewise possible to form at least a part of the wall or walls of the access member of a net material of eg. metal.

The access member may comprise one wall forming a substantially hose-shaped access member, which provides for a simple manufacture of the access member. The access member may eg. be produced by extrusion or by any other method which provides a preferably seam-less access member.

Alternatively, the access member may comprise at least two walls which are formed by sheets of material having substantially larger dimensions in the longitudinal direction than in the transverse direction and being joined at the respective longitudinally extending edges. By this design, a particularly effective sealing is provided. The sheets may eg. be joined by means of welding, adhesion or any other suitable joining technique.

10 In order to control the insertion of the access member properly, said sheets may have different thicknesses and different degrees of flexibility. Hereby, it is possible to control the rigidity in the axial direction of the access member.

15 In an embodiment, which is particularly advantageous with respect to the insertion, at least one blind hole is provided in at least one of said sheets.

In a further embodiment, in which there are at least three sheets and two cavities, and which is particularly advantageous with respect to the insertion as well, one of said cavities is closed at a distance from the outer end of the access member.

In both of these latter embodiments, a suitable fluid, eg. air, may be introduced into the blind hole alternatively the closed cavity, thus increasing the rigidity of the access member in the longitudinal direction thereof during insertion of a catheter whereby the insertion is facilitated.

30 In an embodiment, which is relatively simple to manufacture and which provides for an easy insertion, the inner end of the access member is designed as a cap having a number of openings.

The access member may furthermore comprise means for securing the outer end of the access member to the

abdominal skin surface. Said means may eg. comprise a plate-shaped member, which may be fastened to the skin surface by means of sewing or by adhesion.

In order to provide for additional security
5 against leakage into the access member from the outside, a plug may be provided for introduction into the outer end of said at least one through-going cavity.

In another aspect of the invention, a system for
10 catheterization is provided.

In yet another aspect, a method of replacing an access member is provided. Replacement of the access member may take place by removing the existing access member and shortly after inserting the new access
15 member. If necessary, the new access member may be introduced through the existing one while still in place, whereafter the old one is removed.

BRIEF DESCRIPTION OF THE DRAWINGS.

20 In the following the invention will be described in detail with reference to the schematic drawings, in which

Fig. 1 shows a side view of a system according to the invention during catheterization;

25 Fig. 2 shows a side view of an access member according to the invention in a position of use;

Fig. 3 shows, at a larger scale, a part sectional view of a detail of a system according to another embodiment of the invention;

30 Figs. 4 to 8 show, at a larger scale and very schematic, cross-sectional views of different embodiments of an access member according to the invention; and

Fig. 9 shows a view corresponding to Fig. 2 of a
35 further embodiment of an access member according to the

invention.

DETAILED DESCRIPTION OF THE INVENTION.

In Figs. 1 and 2 a system for suprapubic catheterization is shown, in which an access member 1 is shown in its position of use in a canal extending from the skin surface 2 of the abdominal wall of the user, which in this case is a female, to the urinary bladder 3, said canal extending above the pubic bone 4.

10 The access member 1 is essentially formed as a hollow tube made from a suitable flexible material. The term "tube" should be interpreted in its broadest sense, ie. as comprising any element having at least one longitudinally extending cavity.

15 The wall or walls of the access member is/are formed with a small thickness, which in this respect means that the thickness should be sufficiently low so as to be able to allow parts of one wall, or different walls to contact each other. A preferred thickness

20 depends on the material chosen.

Examples of suitable materials are eg. film or foil made from polyethylene, polyurethane, polypropylene or like material, a flexible foam made from any suitable material, artificial blood vessels, pig

25 guts, Tripsin, a gel, such as a hydrogel or a silicone gel which are widely used for eg. implants or any other gel, or any other material which can meet the demands to the access member, both with respect to physical properties and bio-compatibility. In addition to being

30 flexible and being able to be produced in a small thickness, the material should thus preferably be soft, possess low surface friction, be able to be coated, welded, heat-sealed and/or glued, adhered or joined using any other suitable joining technique and be

35 hydrophobic. Furthermore, the material should be able

to collapse in a radial direction but preferably be stable axially, and could for insertion purposes be rolled up. With respect to the bio-compatibility of the material, it should prevent stenosis, encrustation and
5 bio-film formation, not form in-growth with tissue and be non-toxic.

Parts of the access member may comprise different materials. For instance, the part situated in the region of the abdominal wall could be designed of a net
10 material of eg. metal.

In order to prevent or reduce even further these unwanted effects, the access member may be provided with a coating on the outer side and/or the inner side. The coating may eg. contain antibacterial agents or
15 disinfectants known per se, such as metal ions, halogen ions, antibiotics or sulpha. It is also possible that the wall or walls of the access member may have properties allowing slow release of any known antibacterial or disinfective substances.

20 The access member 1 has an outer end 1a which may be secured to the skin surface 2 by any suitable means, eg. a medical grade adhesive, and an inner end 1b which protrudes well into the bladder 3, the outer and inner ends 1a, 1b defining a predetermined length. Examples of
25 suitable adhesives are adhesives based on styrene-isoprene-styrene block polymer (SIS), polyisobutylene (PIB), Silicone Tacky Gel, polyvinylether (PVE) and acrylic polymers. In the embodiment shown, the cavity in the access member 1 extends throughout the
30 predetermined length such that a catheter 5 may be inserted through the canal provided by the access member 1 in order to attain the catheterization position as shown in Fig. 1, in which urine flows from the bladder 3 through inlet openings 5a provided at the
35 end of the catheter and out to a suitable draining

means (not shown).

After catheterization, the catheter 5 is retracted from the bladder 3 through the access member 1 which remains seated in the body of the user.

5 As indicated in Fig. 2 the access member 1 assumes, at least partially, a flattened position between catheterizations as a result of the involuntary contraction of the detrusor and abdominal muscles, and of the pressure exerted by the urine collected in the
10 bladder, respectively. Consequently, the passage between the bladder 3 and the outside of the body provided by the cavity in the access member is kept closed such that virtually no urine may penetrate to the outside. Moreover, the closure of the canal implies
15 that liquid such as water will not seep into the bladder when the user for example washes, takes a shower or bathes.

Initial positioning of the access member 1 may take place by first penetrating the abdominal wall and
20 the wall of the bladder 3 by means of a trocar and by subsequently inserting a catheter or other applicator means carrying on its outer or inner side the access member 1.

In order to insert the access member 1 without
25 discomfort to the user, the exterior surface of the access member may be provided with a coating to provide a slippery low-friction surface character. In order to retain the access member safely within the body the coating may be of a temporary character such that the
30 exterior surface after a predetermined period of time loses its low-friction character.

Alternatively, application of the access member may take place as shown in Fig. 3, showing a part of an embodiment of the inventive system comprising a
35 catheter 25 and an access member 21. In this

embodiment, an inner end 21b of the access member 21 adapted to be positioned at the end of the catheter 25 provided with urine inlet openings 25a is designed as a cap having openings 21c which allow urine to flow into the catheter 25 through the inlet openings 25a.

In the following, different designs of the access member will be described with reference to Figs. 4 to 8. In these very schematic cross-sectional views, certain details of the access member may be omitted, ie. the access member may comprise parts not indicated in these Figures.

In its most simplified form as shown in Fig. 4, the access member 41 comprises only one circumferential wall 42 which defines a cavity 45 for receiving a catheter during catheterization, thus providing the access member 41 with a substantially hose-shaped appearance. It should be noted that the access member 41 is shown in an open or catheter-receiving position, and it is to be understood that the cavity 45 is kept closed between catheterizations as parts of the wall 42 are pressed against each other.

In the Fig. 5 embodiment, the access member 51 comprises two walls which are formed by sheets 52, 53 of material having substantially larger dimensions in the longitudinal direction than in the transverse direction and being joined at the respective longitudinally extending edges. The cavity 55 defined by the sheets 52, 53 is shown in a slightly open position for reasons of clarity only. In one sheet 53, a blind hole 54 is provided in any suitable manner. During insertion of the access member 51 into the canal, a suitable fluid, eg. air, is introduced into the blind hole 54. As long as the fluid is present in the hole 54, the rigidity of the access member in the longitudinal direction thereof is increased, and the insertion of the access member 51

into the canal is eased.

A similar principle is shown in Fig. 6, in which the access member 61 comprises three walls likewise formed by sheets 62,63,64 of any suitable material, of which sheets 62 and 63 define the catheter receiving cavity 65. The cavity 66 defined between sheets 64 and 63 is closed at a distance from the outer end of the access member, and eg. air may be introduced into the closed cavity 66 in order to ease insertion of the access member 61.

In the Fig. 7 embodiment, the sheets 72,73 forming the walls of the access member 71 have different thicknesses and may in addition thereto have different degrees of flexibility. In this manner secure closing of the cavity as well as an eased introduction is ensured. In addition or alternatively, the thickness and/or the degree of flexibility may vary in the circumferential direction of the access member.

In the embodiment shown in Fig. 8 the cavity 85 defined by the sheets 82,83 forming the walls of the access member 81 is filled with a gel 86, which functions partly as a lubricant during insertion of the catheter, partly as an additional security against leakage.

In Fig. 9 an access member 91 which may be of any of the types described in the above is at its outer end 91a fastened to a plate-shaped member 93, eg. by means of a layer of adhesive 92 or in any other way, such as eg. by forming the plate-shaped member 93 integrally with the access member 91. The plate-shaped member 93 is in turn fastened to the abdominal skin surface by means of eg. a layer 94 of medical grade adhesive. A plug member 95 which is intended to be inserted into the outer end 91a of the access member 91 provides for increased safety against in-seeping of eg. water into

the access member 91. The plug member 95 may be coated as described in the above in connection with the coating of the access member itself.

The access member and the system according to the invention may alternatively be used in urethral catheterization. By using an access member in connection with urethral catheterization, self-catheterization may be performed even by users having a reduced dexterity and mobility as an access member facilitates the operation of finding the urethral orifice, especially in women. In contrast to permanent catheterization the muscles are furthermore allowed to contract and relax. By letting the outer end protrude from the urethral orifice, this end may easily be gripped by the user in order to position the catheter correctly. This operation is thus much facilitated in relation to urethral catheterization without an access member and makes it possible for even eg. sclerosis patients to perform intermittent self-catheterization which in turn implies that this group of patients gains a significantly improved quality of life in relation to use of permanent catheterization.

The access member or system may likewise be used for introduction of eg. pharmaceuticals into the urinary bladder or for washing/rinsing the bladder.

The invention is not limited to the embodiments shown and described in the above. Several modifications and combinations of the embodiments shown and described are conceivable within the scope of the appended claims.

C L A I M S

1. An access member adapted to be, in a position of use, accommodated in an artificial or a natural canal in a user, said access member having an outer end and an inner end and extending, in the position of use, from the outside of the body of the user through said canal and into the urinary bladder, and comprising at least one wall defining at least one cavity extending substantially throughout said predetermined length, said at least one cavity being intended for intermittently receiving a catheter, characterized in that said at least one wall of the access member has such a degree of flexibility that said at least one cavity is kept in a substantially closed position by the mutual contact of parts of said at least one wall, but allows for intermittent insertion of a catheter.

2. An access member according to claim 1, characterized in that the wall or walls of the access member comprise(s) a foil or film material.

3. An access member according to claim 1, characterized in that the wall or walls of the access member comprise(s) a foam or a gel.

4. An access member according to any of claims 1 to 3, characterized in that at least one part of the wall or walls of the access member comprises a net material of eg. metal.

5. An access member according to any of claims 1 to 4, characterized by comprising one wall forming a substantially hose-shaped access member.

6. An access member according to any of claims 1 to 4, characterized by comprising at least two walls which are formed by sheets of material having substantially larger dimensions in the longitudinal direction than in the transverse direction and being

joined at the respective longitudinally extending edges.

7. An access member according to claim 6, characterized in that said sheets are joined by means of welding, adhesion or any other suitable joining technique.

8. An access member according to claim 6 or 7, characterized in that said sheets have different thicknesses.

9. An access member according to any of claims 6 to 8, characterized in that said sheets have different degrees of flexibility.

10. An access member according to any of claims 6 to 9, characterized in that at least one blind hole is provided in at least one of said sheets.

11. An access member according to any of claims 6 to 9, in which there are at least three sheets and two cavities, characterized in that one of said cavities is closed at a distance from the outer end of the access member.

12. An access member according to any of the preceding claims, characterized in that the inner end of the access member is designed as a cap having a number of openings.

13. An access member according to any of the preceding claims, characterized by comprising means for securing the outer end of the access member to the abdominal skin surface.

14. An access member according to claim 13, characterized in that said means comprises a plate-shaped member.

15. An access member according to claim 13, characterized in that the plate-shaped member is fastened to the skin surface by means of an adhesive.

16

16. An access member according to any of the preceding claims, characterized in that a plug member is provided for insertion into the outer end of said at least one through-going cavity.

5 17. A system for catheterization of the urinary bladder through an artificial or a natural canal in a user, comprising a catheter adapted to be inserted through the canal, and an access member according to any of claims 1 to 16.

10 18. A method of replacing an access member according to claim 1, in which a first access member positioned in said canal is removed and a second, substitute access member is inserted shortly afterwards, or a second, substitute access member is
15 introduced through the first access member positioned in said canal whereafter the first access member is removed.

1001945.010802

Table 1

Reaction	Time (min)	Yield (%)	Melting point (°C)	IR (cm ⁻¹)	NMR (ppm)
1	0	0	—	—	—
2	10	10	—	—	—
3	20	20	—	—	—
4	30	30	—	—	—
5	40	40	—	—	—
6	50	50	—	—	—
7	60	60	—	—	—
8	70	70	—	—	—
9	80	80	—	—	—
10	90	90	—	—	—
11	100	100	—	—	—
12	110	110	—	—	—
13	120	120	—	—	—
14	130	130	—	—	—
15	140	140	—	—	—
16	150	150	—	—	—
17	160	160	—	—	—
18	170	170	—	—	—
19	180	180	—	—	—
20	190	190	—	—	—
21	200	200	—	—	—
22	210	210	—	—	—
23	220	220	—	—	—
24	230	230	—	—	—
25	240	240	—	—	—
26	250	250	—	—	—
27	260	260	—	—	—
28	270	270	—	—	—
29	280	280	—	—	—
30	290	290	—	—	—
31	300	300	—	—	—
32	310	310	—	—	—
33	320	320	—	—	—
34	330	330	—	—	—
35	340	340	—	—	—
36	350	350	—	—	—
37	360	360	—	—	—
38	370	370	—	—	—
39	380	380	—	—	—
40	390	390	—	—	—
41	400	400	—	—	—
42	410	410	—	—	—
43	420	420	—	—	—
44	430	430	—	—	—
45	440	440	—	—	—
46	450	450	—	—	—
47	460	460	—	—	—
48	470	470	—	—	—
49	480	480	—	—	—
50	490	490	—	—	—
51	500	500	—	—	—
52	510	510	—	—	—
53	520	520	—	—	—
54	530	530	—	—	—
55	540	540	—	—	—
56	550	550	—	—	—
57	560	560	—	—	—
58	570	570	—	—	—
59	580	580	—	—	—
60	590	590	—	—	—
61	600	600	—	—	—
62	610	610	—	—	—
63	620	620	—	—	—
64	630	630	—	—	—
65	640	640	—	—	—
66	650	650	—	—	—
67	660	660	—	—	—
68	670	670	—	—	—
69	680	680	—	—	—
70	690	690	—	—	—
71	700	700	—	—	—
72	710	710	—	—	—
73	720	720	—	—	—
74	730	730	—	—	—
75	740	740	—	—	—
76	750	750	—	—	—
77	760	760	—	—	—
78	770	770	—	—	—
79	780	780	—	—	—
80	790	790	—	—	—
81	800	800	—	—	—
82	810	810	—	—	—
83	820	820	—	—	—
84	830	830	—	—	—
85	840	840	—	—	—
86	850	850	—	—	—
87	860	860	—	—	—
88	870	870	—	—	—
89	880	880	—	—	—
90	890	890	—	—	

1. The first part of the document is a title page. It contains the title "THE HISTORY OF THE UNITED STATES OF AMERICA" and the author "BY JAMES M. SMITH".

2. The second part of the document is a table of contents. It lists the chapters and their corresponding page numbers.

3. The third part of the document is the first chapter, titled "THE DISCOVERY OF AMERICA". It describes the early exploration of the continent by Christopher Columbus and other European navigators.

4. The fourth part of the document is the second chapter, titled "THE SETTLEMENT OF AMERICA". It discusses the early colonial settlements and the challenges faced by the settlers.

5. The fifth part of the document is the third chapter, titled "THE REVOLUTIONARY WAR". It covers the events leading up to the war and the battle of independence.

6. The sixth part of the document is the fourth chapter, titled "THE CONSTITUTION". It explains the formation of the federal government and the principles of the Constitution.

7. The seventh part of the document is the fifth chapter, titled "THE WESTERN EXPANSION". It describes the westward movement of the population and the acquisition of new territories.

8. The eighth part of the document is the sixth chapter, titled "THE CIVIL WAR". It details the conflict between the Union and the Confederacy and its impact on the nation.

9. The ninth part of the document is the seventh chapter, titled "THE RECONSTRUCTION". It discusses the efforts to rebuild the South and the challenges of integrating freed slaves into society.

10. The tenth part of the document is the eighth chapter, titled "THE MODERN UNITED STATES". It covers the period from the end of the Civil War to the present day, focusing on economic growth and social change.

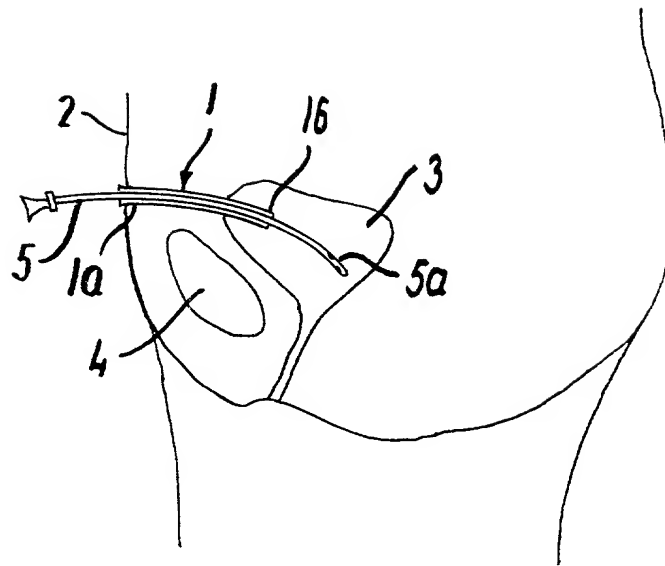


FIG. 1

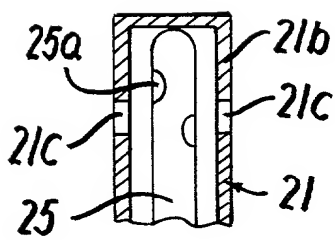


FIG. 3

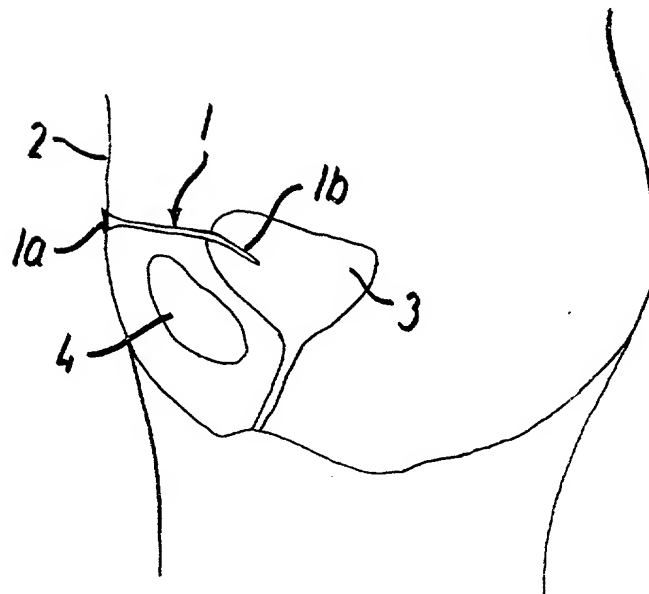


FIG. 2

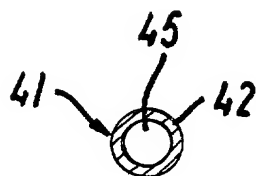


FIG. 4

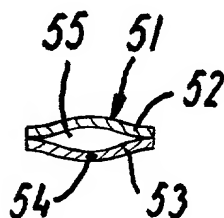


FIG. 5

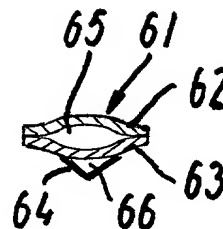


FIG. 6

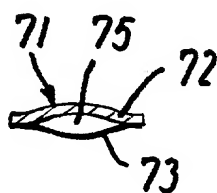


FIG. 7

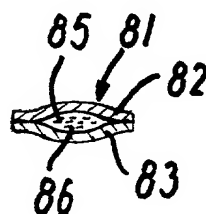


FIG. 8

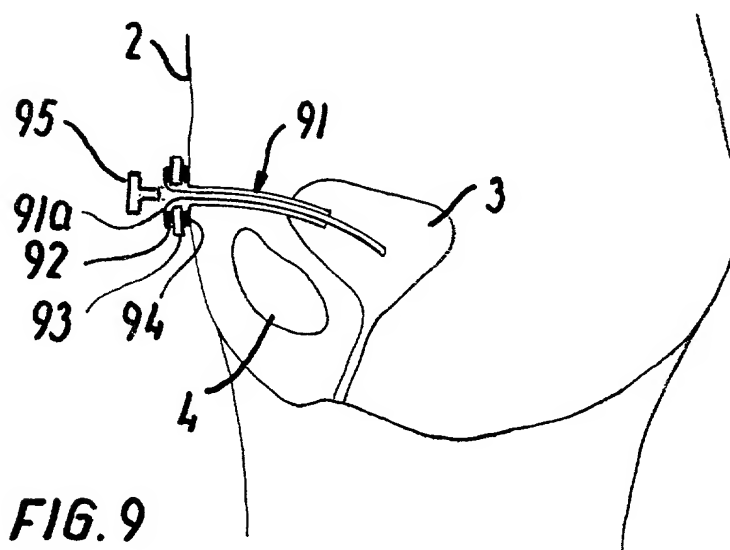


FIG. 9

DECLARATION
AND POWER OF ATTORNEY
U.S.A.

FOR ATTORNEYS' USE ONLY

ATTORNEYS' DOCKET NO.

ALL PATENTS, INCLUDING DESIGN

FOR APPLICATION BASED ON PCT; PARIS CONVENTION;

NON PRIORITY; OR PROVISIONAL APPLICATIONS

As a below named inventor, I declare that my residence, post office address and citizenship are stated below next to my name, the information given herein is true, that I believe that I am the original, first and sole inventor (if only one name is listed at 201 below), or an original, first and joint inventor (if plural inventors are named below at 201-203, or on additional sheets attached hereto) of the subject matter which is claimed and for which patent is sought on the invention entitled: "AN ACCESS MEMBER AND A SYSTEM FOR CATHETERIZATION OF THE URINARY BLADDER THROUGH AN ARTIFICIAL OR A NATURAL CANAL IN A USER, AND A METHOD OF REPLACING SUCH AN ACCESS MEMBER"

which is described and claimed in: ☒ PCT International Application No. PCT/DK00/00361 filed 03 July 2000
☐ the attached specification ☐ the specification in application Serial No. _____ filed _____
(if applicable) and amended on 20 September 2001

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, §119 (a)-(d) of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

Prior Foreign Application(s)

PA 1999 00958
(Number)

Denmark
(Country)

02 July 1999
(Day/Month/Year Filed)

Priority Claimed

☒ Yes ☐ No

(Number)

(Country)

(Day/Month/Year Filed)

☐ Yes ☐ No

(Number)

(Country)

(Day/Month/Year Filed)

☐ Yes ☐ No

I hereby claim the benefit under Title 35, United States Code, §119(e) of any United States provisional application(s) listed below:

Application No. _____ Filing Date _____ Application No. _____ Filing Date _____

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application:

(Application Serial No.)

(Filing Date)

(Status: patented, pending, abandoned)

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorneys (Registration No.) to prosecute this application, receive and act on instructions from my agent, and transact all business in the Patent and Trademark Office connected therewith. HARVEY B. JACOBSON, JR. (20,851); D. DOUGLAS PRICE (24,514); JOHN CLARKE HOLMAN (22,769); MARVIN R. STERN (20,640); MICHAEL R. SLOBASKY (26,421); JONATHAN L. SCHERER (29, 851); IRWIN M. AISENBERG (19,007); WILLIAM E. PLAYER (31,409)

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PROFESSIONAL LIMITED LIABILITY COMPANY

*Inventor(s) name must include at least one unabbreviated first or middle name.

201	FULL NAME* OF INVENTOR	FAMILY NAME	GIVEN NAME	MIDDLE NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE OR COUNTRY
202	FULL NAME* OF INVENTOR	FAMILY NAME	GIVEN NAME	MIDDLE NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE OR COUNTRY
203	FULL NAME* OF INVENTOR	FAMILY NAME	GIVEN NAME	MIDDLE NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE OR COUNTRY

I further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment or both, under section 1001 of Title 18 of the United States Code; and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

SIGNATURE OF INVENTOR 201*	SIGNATURE OF INVENTOR 202*	SIGNATURE OF INVENTOR 203*
DATE	DATE	DATE

☐ Additional inventors are named on separately numbered sheets attached hereto.

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